Intracellular Immunization of Rhesus CD34⁺ Hematopoietic Progenitor Cells With a Hairpin Ribozyme Protects T Cells and Macrophages From Simian Immunodeficiency Virus Infection

By Michael Rosenzweig, Douglas F. Marks, Donna Hempel, Marina Heusch, Gunter Kraus, Flossie Wong-Staal, and R. Paul Johnson

Evaluation of candidate genes for stem cell gene therapy for acquired immunodeficiency syndrome (AIDS) has been limited by the difficulty of supporting in vitro T-cell differentiation of genetically modified hematopoietic progenitor cells. Using a novel thymic stromal culture technique, we evaluated the ability of a hairpin ribozyme specific for simian immunodeficiency virus (SIV) and human immunodeficiency virus type 2 (HIV-2) to inhibit viral replication in T lymphocytes derived from transduced CD34⁺ progenitor cells. Retroviral transduction of rhesus macaque CD34⁺ progenitor cells with a retroviral vector (p9456t) encoding the SIV-specific ribozyme and the selectable marker neomycin phosphotransferase in the presence of bone marrow stroma and in the absence of exogenous cytokines resulted in efficient transduction of both colony-forming units and long-term culture-initiating cells, with transduction efficiencies ranging between 21% and 56%. After transduction, CD34⁺ cells were cultured on rhesus thymic stromal culture (to support in vitro differentiation of T cells) or in the presence of cytokines (to support differentiation of macrophage-like cells). After expansion and selection with the neomycin analog G418, cells derived from transduced progenitor cells were challenged with SIV. CD4⁺ T cells derived from CD34⁺ hematopoietic cells transduced with the ribozyme vector p9456t were highly resistant to challenge with SIV, exhibiting up to a 500-fold decrease in SIV replication, even after high multiplicities of infection. Macrophages derived from CD34⁺ cells transduced with the 9456 ribozyme exhibited a comparable level of inhibition of SIV replication. These results show that a hairpin ribozyme introduced into CD34⁺ hematopoietic progenitor cells can retain the ability to inhibit AIDS virus replication after T-cell differentiation and support the feasibility of intracellular immunization of hematopoietic stem cells against infection with HIV and SIV. Protection of multiple hematopoietic lineages with the SIV-specific ribozyme should permit analysis of stem cell gene therapy for AIDS in the SIV/macaque model.

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genetically modified human CD34+ hematopoietic progenitor cells. In this study, CD34+ cells were transduced with a retroviral vector encoding the hairpin ribozyme MJT, which cleaves a conserved sequence contained in the HIV-1 LTR. Similar observations have been recently reported by others who have shown inhibition of HIV-1 replication in myelomonocytic cells derived from CD34+ cells transduced with retroviral vectors encoding a rev-responsive element decoy, a transdominant Rev protein, or a double hammerhead ribozyme. However, until recently, no data were available on whether genes designed to inhibit HIV-1 or simian immunodeficiency virus (SIV) replication could be introduced into CD34+ progenitor cells and remain active after T-cell differentiation. The lack of information on this critical issue arises in part from the fact that T-cell differentiation normally occurs in the unique three-dimensional environment of the thymus and has been difficult to model using in vitro culture systems.

Because properties of genes designed to inhibit HIV-1/ SIV replication may differ between macrophages and T cells, both with respect to expression, toxicity, and efficacy, it is important to analyze the ability of candidate therapeutic genes for stem cell therapy for AIDS for their ability to inhibit viral replication in T cells as well as macrophages derived from transduced progenitor cells. In the present study, we evaluate the ability of an SIV-specific hairpin ribozyme to inhibit viral replication in T cells and macrophages derived from transduced CD34+ progenitor cells. Infection of macaques with SIV is generally considered to be CD3, CD4, and CD8. After staining, cells were sorted into CD3+CD4+CD8+ and CD3+CD4+CD8- subsets using a Becton Dickinson FACS Vantage. Fluorescence was excited using an argon Coherent Enterprise laser (Mountain View, CA) generating 90 mW of 488 nm light. Fluorocin isothiocyanate emission was detected through a 530/30-nm bandpass filter and phycoerythrin emission was detected through a 575/26-nm bandpass filter. Fluorescence was directed to the appropriate detectors using a 560-nm short pass dichroic.

**MATERIALS AND METHODS**

**Bone marrow harvest and immunomagnetic bead CD34 cell isolation.** Rhesus monkeys (Macaca mulatta) used in this study were normal, colony-born animals maintained in accordance with the guidelines of the Committee on Animals of the Harvard Medical School and the Guide for the Care and Use of Laboratory Animals (DHHS Publication No. [NIH] 85-23, revised 1985). Bone marrow aspirates were obtained from anesthetized macaques and CD34+ cells purified by immunomagnetic beads (Dynal, Lake Success, NY), as previously described. Rhesus CD34+ cells obtained in this fashion were greater than 90% CD34+ and contained less than 1% residual CD3- T cells. Purified CD34+ cells were subsequently further depleted of CD3- cells by incubation with a CD3-specific antibody 6G12 (kindly provided by J. Weng, Massachusetts General Hospital, Boston, MA), followed by incubation with sheep antimouse magnetic beads (Dynal) and depletion of cells bound to beads using a magnetic separation device (Dynal).

**Flow cytometric analysis and cell sorting.** Antibodies used for immunophenotyping of rhesus cells included anti-CD3 (6G12), anti-CD4 (OKT4; obtained from the American Type Culture Collection, Rockville, MD), anti-CD8 (Leu-2a; Becton Dickinson, San Jose, CA), anti-CD14 (MOP9; Becton Dickinson), and anti-CD34 (QBend-10; Immunotech, Westbrook, ME). Cells were stained in the presence of phosphate-buffered saline with 2% mouse serum. Cells were harvested from in vitro T-cell cultures and incubated with antibodies to CD3, CD4, and CD8. After staining, cells were sorted into CD3+CD4+CD8- and CD3+CD4+CD8+ subsets using a Becton Dickinson FACSort Vantage. Fluorescence was excited using an argon Coherent Enterprise laser (Mountain View, CA) generating 90 mW of 488 nm light. Fluorescein isothiocyanate emission was detected through a 530/30-nm bandpass filter and phycoerythrin emission was detected through a 575/26-nm bandpass filter. Fluorescence was directed to the appropriate detectors using a 560-nm short pass dichroic.

**Retroviral vectors and transduction.** Retroviral vectors used in these studies are diagrammed in Fig 1. The vector p9456t encodes an SIV-specific hairpin ribozyme 9456t expressed under the control of a rRNA promoter in the retroviral vector LNL6. Two control retroviral vectors were used: MJT, which encodes an HIV-1--specific ribozyme that does not recognize SIV, and the unmodified vector LNL6. All retroviral vectors were produced in the murine amphototropic packaging cell line PA317. Retroviral packaging cell lines were grown in an artificial capillary system (ACS; Cellco, Germantown, MD). All transductions were performed using frozen supernatants with titers ranging from 1.0 to 1.6 × 106 colony-forming units (CFU)/mL as determined by end point dilution on 3T3 cells. No replication competent retrovirus was detected in retroviral supernatants as assessed using the S′L- assay employing PG4 cells. Rhesus CD34+ bone marrow cells were transduced for 3 days with cell-free retroviral vector supernatants. Transductions were performed in the presence of irradiated allogeneic bone marrow stroma. Rhesus bone marrow stromal cultures were established in long-term culture media (Stem Cell Technologies, Vancouver, British Columbia, Canada) plus 10⁻³ mol/L hydrocortisone (Sigma, St Louis, MO) and γ irradiated (15 Gy) after 7 to 14 days of culture before replating. Nonadherent cells were removed after 3 days of transduction, counted, and used to establish in vitro T lymphopoiesis or macrophage cultures as described below.

**T-cell differentiation of CD34+ bone marrow cells.** To support differentiation into T cells, transduced CD34+ cells were cultured...
on a thymic stromal monolayer as previously described. Briefly, thymic stromal monolayers were established in 24-well plates from cryopreserved fetal rhesus thymus cell suspensions and nonadherent cells were vigorously washed away 3 days after. After 7 days, thymic stromal monolayers were inoculated with 2 x 10^7 CD34^+ cells/well and cultured in the presence of RPMI with 10% fetal calf serum (FCS). After 14 days, samples were removed for flow cytometry and polymerase chain reaction (PCR) analysis, and the remaining cells were expanded by stimulation with Concanavalin A (ConA; 5 μg/mL; Gibco, Cambridge, MA) and irradiated human peripheral blood mononuclear cells (PBMC).

Aliquots of T cells were analyzed by three-color flow cytometry and CD3^+ CD4^- CD8^- cells were obtained by cell sorting as described above. Purified CD4^+ T cells were then restimulated with ConA, IL-2, and irradiated PBMC and selected in G418 (200 μg/mL, active concentration; Gibco BRL, Gaithersburg, MD) for 7 days before restimulation and SIV infection.

** Colony-forming assays and differentiation of macrophage-like cells from CD34^+ cells.** CFU assays were established by culturing CD34^+ cells at a concentration of 1 x 10^4 cells/mL in complete methylcellulose media (Stem Cell Technologies) containing human erythropoietin (10 ng/mL; Stem Cell Technologies), granulocyte-macrophage colony-stimulating factor (GM-CSF; 30 ng/mL; Genzyme, Cambridge, MA), human IL-3 (20 ng/mL; Genzyme), and human stem cell factor (SCF; 50 ng/mL; Genzyme) in the presence and absence of G418 (400 μg/mL, active concentration; Gibco BRL). Hematopoietic colonies were scored at 14 and 21 days based on standard criteria. Transduction efficiency was calculated by dividing the number of G418-resistant CFU by the total number of CFU for each culture condition examined. Previous experiments have shown that greater than 95% of G418-resistant CFU obtained under these conditions contain neo vector DNA. Individual myelomonocytic colonies were pooled, counted, and cultured in 24-well plates in Dulbecco's modified Eagle's medium with 10% fetal bovine serum and supplemented with GM-CSF (20 ng/mL) and macrophage colony-stimulating factor (M-CSF; 40 U/mL; Genzyme) to induce differentiation of macrophage-like cells. After 7 days of selection, the resulting cells were infected with SIV as described below.

**Proliferation assays.** T cells derived from CD34^+ cells were washed and resuspended in RPMI with 10% FCS at a concentration of 1 x 10^5 cells/mL. One hundred microliters (10^5 cells) were added to each well of a 96-well plate. Cells were stimulated with either ConA (5 μg/mL) or monoclonal antibodies to CD34 (6G12) and CD28 (2.2; ImmunoTech, Larrabee, ME) (0.15 μg/mL) in the presence of IL-2 (20 U/mL) and irradiated human PBMC (10^5 cells/well in 100 μL of RPMI with 10% FCS). Purified goat antimouse F(ab')2 fragments (Kirkegard and Perry Laboratories, Gaithersburg, MD) were used as a cross-linking agent for the stimulation with the monoclonal antibodies to CD3 and CD28. Wells were pretreated with 1.25 μg/mL of goat antimouse antibody for 45 minutes at 37°C and washed three times before the addition of monoclonal antibodies to CD3 and CD28. After 7 days in culture at 37°C, each well was pulsed with 1 μCi of methyl-H-thymidine (DuPont, NEN, Boston, MA) for 16 hours and harvested onto a glass fiber filter (Harvester 96; Tomtec, Orange, CT) and the incorporation of H-thymidine was determined using a liquid scintillation counter (Micro Beta Plus; Wallac, Gaithersburg, MD). Five replicate wells were assayed for each condition.

**PCR analysis of retroviral genes in T cells.** Cellular DNA was extracted using the Puregene DNA isolation kit (Gentra Systems, Inc, Minneapolis, MN). DNA template (100 ng, corresponding to approximately 10^7 cell equivalents) was added to each PCR reaction (50 μL) containing 50 ng each of the sense and antisense PCR primer, 1.0 mmol/L MgCl2, 200 μmol/L of each dNTP, and 2.5 U of AmpliTaq (Perkin Elmer, Norwalk, CT). The neo primer (5' primer, listed 5' to 3') TCC ATC ATG GCT GAT GCA TGG C and (3' primer) GAT AGA AGG CGA TGC GCT GGC AAT CG yield a 416-bp product; the 5' PCR primer was end-labeled with 32P-dATP (DuPont, NEN). Amplification was performed at 92°C for 2 minutes, followed by 35 cycles of 92°C for 30 seconds, 65°C for 30 seconds, and 72°C for 1 minute and a final extension at 72°C for 10 minutes using a PTC 200 DNA Engine (MJ. Research, Watertown, MA). To facilitate semiquantitative analysis of samples, each PCR amplification included amplification of a series of standards consisting of 10-fold dilutions (100%, 10%, 1%, and 0.1%) of the CEMx174 cell line stably transduced with the LN retroviral vector diluted in genomic rhesus DNA. PCR products were separated on a 4% native polyacrylamide gel and then visualized by autoradiography using Biomax MR Film (Eastman Kodak, Rochester, NY).

**SIV infection of T cells and macrophage-like cells.** Rhesus CD4^+ T-cell progeny and macrophage-like cells derived from CD34^- cells transduced with the ribosome or the control vectors were infected with SIVmac239 (T cells) or SIVmac316 (macrophage-like cells) for 4 hours at 37°C. SIVmac239 is a pathogenic molecular clone of SIV that replicates efficiently in T cells. SIVmac316 is derived from SIVmac239, differing in 8 amino acids in the envelope sequence, and replicates well in macrophages. T cells were challenged with SIVmac239 at multiplicities of infection (MOI) ranging from 0.001 to 1.0 TCID50/cell; macrophage-like cells were challenged with SIVmac316 at MOI of 0.001 to 0.01 TCID50/cell. The titer of virus stocks was performed by endpoint dilution on CEMx174 cells performed in 6 replicate wells as previously described for HIV-1. Free virus was removed by vigorously washing three times with phosphate-buffered saline. Infections were performed in duplicate and the results were presented as the mean of each well. Viral replication was assessed by enzyme-linked immunosorbent assay determination of SIV p27 antigen (Coulter, Hialeah, FL) according to the manufacturer’s protocol. The lower limit detection for SIV p27 is 50 pg/mL.

**RESULTS**

Transduction of rhesus CD34^+ progenitor cells with retroviral vectors encoding the SIV-specific ribozyme 9456. We first evaluated retroviral transduction efficiency of CD34^- progenitor cells as assessed by outgrowth of CFU resistant to the neomycin analog G418. G418-resistant erythroid and myelomonocytic CFU were assessed after transduction with a retroviral vector encoding an SIV-specific ribozyme (p9456t) and the neomycin phosphotransferase gene neo or the LNL6 vector encoding neo alone. Based on the observation that stimulation with exogenous cytokines inhibits the ability of CD34^- cells to undergo T-cell differentiation (Rosenzweig et al, manuscript in preparation), CD34^- cells were transduced by three successive incubations with cell-free retroviral supernatant in the presence of bone marrow stroma and the absence of exogenous cytokines. Using this approach, transduction efficiencies of 21% to 35% were obtained using the p9456t vector and 30% to 45% using the control LNL6 vector (Table 1). Similar numbers of myelomonocytic (colony-forming unit–granulocyte-macrophage [CFU-GM]) and erythroid (burst-forming unit-erythroid [BFU-E]) colonies were obtained after transduction with the p9456t and LNL6 vectors, suggesting that 9456 ribozyme did not interfere with differentiation of myelomonocytic or erythroid lineages.

To assess transduction of more primitive hematopoietic...
cells, we also analyzed the yield of G418-resistant long-term culture-initiating cells (LTC-IC) obtained from transduced CD34⁺ cells. Rhesus CD34⁺ progenitor cells were transduced as described above, cultured on irradiated bone marrow stroma for 5 weeks, and then replated in methylcellulose in the presence and absence of G418. After transduction with the p9456t retroviral vector, we observed between 30% and 56% transduction of LTC-IC, and a similar transduction efficiency with the LNL6 vector (Table 2).

Table 1. Transduction Efficiency of Rhesus CD34⁺ Cells by the LNL6 or 9456 Ribozyme Retroviral Vectors

<table>
<thead>
<tr>
<th>Experiment No.</th>
<th>LNL6</th>
<th>p9456t</th>
<th>Untransduced</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>G418⁺ CFU⁺</td>
<td>%G418⁺</td>
<td>M:E</td>
</tr>
<tr>
<td>1</td>
<td>48/150</td>
<td>33%</td>
<td>2:1</td>
</tr>
<tr>
<td>2</td>
<td>90/205</td>
<td>45%</td>
<td>3.2:1</td>
</tr>
<tr>
<td>3</td>
<td>54/180</td>
<td>30%</td>
<td>2:1</td>
</tr>
</tbody>
</table>

* G418-resistant (G418⁺) CFU were determined after culture in methylcellulose with 400 µg/mL G418 (active concentration).
† M:E is the myelomonocytic:erythroid ratio. M:E ratios for CFU derived from transduced CD34⁺ cells were calculated for G418⁺ colonies.

Table 2. Transduction Efficiency of LTC-IC by the LNL6 or 9456 Ribozyme Retroviral Vectors

<table>
<thead>
<tr>
<th>Experiment No.</th>
<th>LNL6</th>
<th>p9456t</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>G418⁺-Resistant CFU⁺</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>35%</td>
<td>37%</td>
</tr>
<tr>
<td>2</td>
<td>54%</td>
<td>56%</td>
</tr>
<tr>
<td>3</td>
<td>33%</td>
<td>30%</td>
</tr>
</tbody>
</table>

* Transduced CD34⁺ cells were cultured on irradiated bone marrow stroma for 5 weeks and then cultured in methylcellulose in the presence and absence of G418. Colony counts reflect the sum of both CFU-GM and BFU-E.
Table 3. Immunophenotyping of T Cells Derived From CD34+ Cells Transduced With the LNL6 or 9456 Ribozyme Retroviral Vectors

<table>
<thead>
<tr>
<th>Day</th>
<th>Stage*</th>
<th>LNL6</th>
<th>p9456t</th>
<th>Untransduced</th>
</tr>
</thead>
<tbody>
<tr>
<td>15</td>
<td>Thymic culture</td>
<td>45</td>
<td>21</td>
<td>34</td>
</tr>
<tr>
<td>28</td>
<td>Expansion</td>
<td>34</td>
<td>18</td>
<td>48</td>
</tr>
<tr>
<td>42</td>
<td>Cell sorting</td>
<td>97</td>
<td>96</td>
<td>87</td>
</tr>
<tr>
<td>56</td>
<td>Expansion</td>
<td>93</td>
<td>96</td>
<td>89</td>
</tr>
<tr>
<td>70</td>
<td>G418</td>
<td>94</td>
<td>94</td>
<td>94</td>
</tr>
<tr>
<td>84</td>
<td>SIV</td>
<td>88</td>
<td>88</td>
<td>88</td>
</tr>
<tr>
<td>98</td>
<td>SIV</td>
<td>96</td>
<td>94</td>
<td>96</td>
</tr>
</tbody>
</table>

Expression of CD4 and CD8 was assessed on CD34+ T cells by three-color flow cytometric analysis. Results are expressed as a percentage of all CD34+ cells.

* CD34+ cells were transduced with the indicated retroviral vector, cultured on thymic stroma for 15 days, and expanded by stimulation with ConA, irradiated human PBMC, and IL-2; then CD4+ and CD8+ T cells were selected by cell sorting. These cells were then restimulated, cultured in the presence of G418 (200 μg/mL G418, active concentration), restimulated, and then infected with SIVmac239. Untransduced CD34+ cells were cultured in parallel up until the point of G418 selection.

discussion

It has been almost 10 years since the strategy of intracellular immunization of CD34+ progenitor cells against AIDS virus infection was first proposed. Since that time, despite multiple reports demonstrating the ability of genes to inhibit HIV or SIV replication in transformed cell lines, there have been relatively few publications that have analyzed inhibition of replication in the progeny of transduced hematopoietic progenitor cells and most of these have analyzed myelomonocytic rather than T cells. In this study, we examined the ability of an SIV-specific hairpin ribozyme to inhibit viral replication in hematopoietic cells derived from transduced CD34+ progenitor cells. Our data show that the 9456 ribozyme is quite effective in inhibiting SIV replication in both T cells and macrophage-like cells derived from trans-
Fig 4. Inhibition of SIV replication in CD4+ T cells derived from CD34+ cells transduced with the 9456 ribozyme. CD34+ cells were transduced with a retroviral vector containing either the 9456 ribozyme (●) or, as controls, the LNL6 vector (■) or the HIV-1-specific ribozyme MJT (▲), which does not recognize SIV. Transduced CD34+ cells were then cultured on thymic stroma to support T-cell differentiation, expanded, selected in G418, and then infected with SIVmac239 at an MOI of 0.01 TCID50 (A), 0.1 TCID50 (B), or 1 TCID50/cell (C).

We observed a greater than 500-fold reduction in viral replication as compared with controls, even after a relatively high MOI of 1 TCID50/cell. In addition, we observed no evident toxic effects on T-cell differentiation, cellular proliferation, or expression of CD4.

Our results add to the expanding body of evidence showing the efficacy of ribozymes for gene therapy and also emphasize their potential value for stem cell gene therapy. The hairpin ribozyme MJT, which cleaves a conserved sequence in the U5 region of the HIV-1 LTR, is able to inhibit replication of HIV-1 in a variety of cell types, including transformed cell lines, primary human T cells, and macrophage-like cells derived from transduced CD34+ bone marrow cells. Efficient expression of ribozymes by RNA polymerase III promoters, which are active in diverse cell types, and the absence of demonstrable toxicity of ribozymes represent distinct advantages of these molecules for stem cell gene therapy, where it is desirable to achieve sustained expression in multiple different hematopoietic lineages without adverse effects. Our results with the SIV-specific 9456 ribozyme suggest that comparable hairpin ribozymes specific for HIV-1 may have a similar capacity to inhibit viral replication in T cells derived from transduced progenitor cells. Possible disadvantages of ribozymes include the potential for the emergence of virus variants with mutations in the target sequence recognized by the ribozyme, a particular concern given the relatively high mutation rate of HIV and the relatively rapid rates of virion turnover in vivo. However, the use of ribozymes in combination with other ribozymes or other gene therapy approaches and the choice of target sequences in highly conserved regions of the viral genome should limit the emergence of resistance.

Several considerations suggest that evaluation of genes designed to inhibit HIV replication in T cells derived from transduced progenitor cells may offer significant information not available from the analysis of myelomonocytic progeny. T cells are the dominant population of HIV-infected cells in vivo and are estimated to account for greater than 99% of viral production on a daily basis. Differentiation of T cells from progenitor cells differs considerably from that of myelomonocytic cells. T-cell differentiation involves a complex series of developmental stages resulting in the rearrangement of T-cell receptor genes and the production of a diverse population of T cells able to recognize foreign antigenic epitopes.

Fig 5. Expression of CD4 and CD14 on macrophage-like cells derived from transduced CD34+ cells. CD34+ cells transduced with either the LNL6 (A) or ribozyme 9456 (B) retroviral vectors were cultured in methylcellulose in the presence of G418 and then replated in liquid culture supplemented with GM-CSF, M-CSF, and G418. Macrophage-like cells derived from these cultures were then analyzed for expression of CD4 and CD14 by flow cytometry.
peptides presented by self major histocompatibility complex molecules. A large number of genes are expressed during the different stages of T-cell development and interference with these genes may adversely influence T-cell production. The potential for toxicity of foreign genes expressed in hematopoietic cells derived from transduced CD34+ progenitor cells may therefore be greater in T cells and their progenitors than in macrophages. In addition, expression of genes in T cells may differ from that in macrophages. Our current data show that introduction of the 9456 ribozyme into CD34+ progenitor cells does not appear to interfere with T-cell differentiation and that the 9456 ribozyme is expressed in mature CD4+ T cells at sufficient levels to inhibit SIV replication. We have also recently shown that a polymeric transactivation response (TAR) element decoy linked to an antisense tat molecule expressed under the control of a modified HIV-1 LTR is able to inhibit both HIV-1 and SIV replication in T cells derived from transduced CD34+ progenitor cells. Similar results have been subsequently reported using a retroviral vector encoding a transdominant mutant HIV-1 RevM10 protein. Taken together, these results demonstrate that candidate therapeutic genes introduced into CD34+ hematopoietic progenitor cells can retain the ability to inhibit AIDS virus replication after T-cell differentiation.

Definition of the optimal conditions for retroviral transduction that maintain the pluripotency of hematopoietic stem cells has been difficult to achieve. Because retroviral vectors based on Moloney murine leukemia virus require cell cycling for integration of vector DNA, most investigators have used stimulation with hematopoietic growth factors to enhance retroviral transduction of hematopoietic progenitor cells. Although cytokine stimulation can significantly enhance retroviral transduction, it may also lead to a loss of pluripotency. Stimulation of murine hematopoietic cells with cytokines significantly decreases their ability to reconstitute irradiated hosts and to differentiate into B cells. We have observed that the cytokines IL-3 and SCF, which are commonly used to enhance retroviral transduction, significantly inhibit the T-cell differentiation capacity of CD34+ progenitor cells (Rosenzweig et al, manuscript in preparation). Because bone marrow stroma has previously been shown to enhance retroviral transduction efficiency and to maintain the long-term engraftment of transduced CD34+ human cells in immunodeficient mice, in these studies we performed retroviral transduction in the presence of bone marrow stroma and the absence of exogenous cytokines. Using these conditions and relatively high titer retroviral supernatant, we were able to obtain efficient transduction of CD34+ cells.

Figure 6. Inhibition of SIV replication in macrophage-like cells derived from CD34+ cells transduced with the 9456 ribozyme. CD34+ cells were transduced with a retroviral vector containing either the 9456 ribozyme (●) or, as controls, the LNL6 vector (■) or the HIV-1 specific ribozyme MJT (▲). After transduction, the macrophage-like cells were derived as described in the legend for Fig 5 and challenged with SIVmac316 using an MOI of 10−2 TCID50/cell.

In vivo studies in macaques should permit examination of the efficiency of retroviral transduction of true hematopoietic stem cells under the conditions used in these experiments. Despite the many potential advantages of stem cell gene therapy for AIDS, there are many obstacles to overcome. The levels of genetically modified hematopoietic cells in most human and nonhuman primate trials of transduced hematopoietic stem cells have been disappointingly low, generally less than 1%. This inefficient level of genetic modification is likely to reflect the difficulties in transducing hematopoietic stem cells, which are largely quiescent, with the current generation of retroviral vectors that are dependent on cell cycling for efficient integration. Additional obstacles to stem cell gene therapy for AIDS are unique to HIV infection. Although CD34+ bone marrow cells appear to be infected only at a very low level (if at all) in vivo and CD34+ cells from HIV-infected subjects can be efficiently transduced with retroviral vectors, there are multiple reports of abnormal hematopoiesis in HIV-infected people (reviewed in Harbol et al). Thus, abnormalities in the bone marrow microenvironment of HIV-infected people might interfere with differentiation of genetically modified hematopoietic stem cells. An additional potential barrier to stem cell gene therapy for AIDS is the loss of thymic function, either due to age or the effects of HIV infection. A final concern is that reservoirs of HIV in relatively long-lived hematopoietic cells (eg, macrophages, dendritic cells, or microglia) or nonhematopoietic cells resident in these microenvironments (eg, thymic epithelial cells) may interfere with the ability of genetically modified cells to differentiate or restore immune function.

In light of these challenges, trials in nonhuman primates are likely to play an important role in the development of
stem cell gene therapy for AIDS. Carefully designed trials in the SIV/macaque model can address the issues of optimization of transduction of hematopoietic stem cells, engraftment and differentiation of genetically modified cells in an SIV-infected host, the potential for evolution of virus resistant to a specific therapeutic gene, and the relative efficacy and toxicity of different approaches.

In sum, our results demonstrate potent inhibition of SIV replication in both T cells and macrophages derived from CD34+ progenitor cells transduced with a hairpin ribozyme. These findings provide strong evidence that intracellular immunization of hematopoietic stem cells with a ribozyme can protect progeny from AIDS virus infection and will permit analysis of stem cell gene therapy approaches for AIDS in nonhuman primates.

ACKNOWLEDGMENT

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Metastatic carcinoid tumor of midgut origin. A 70-year-old woman who had an ileocolectomy for carcinoid tumor 2 months earlier was found to have a pelvic mass. At surgery, bilateral tumors in the ovaries were present and metastases were evident in the liver. The metastatic ovarian tumor shown here contained numerous completely electron-dense, pleomorphic secretory granules diagnostic for carcinoid tumors of midgut origin, in this case from the previously resected ileum. Immunoperoxidase stains were positive for chromogranin and synaptophysin. Original magnification × 8,000 (Courtesy of Ann M. Dvorak, MD, Department of Pathology, Beth Israel Deaconess Medical Center, 330 Brookline Ave, Boston, MA 02215.)
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